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PIG-W Is Critical for Inositol Acylation but Not for Flipping of Glycosylphosphatidylinositol-Anchor

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Many cell surface proteins are anchored to a membrane via a glycosylphosphatidylinositol (GPI), which is attached to the C termini in the endoplasmic reticulum. The inositol ring of phosphatidylinositol is acylated during biosynthesis of GPI. In mammalian cells, the acyl chain is added to glucosaminyl phosphatidylinositol at the third step in the GPI biosynthetic pathway and then is usually removed soon after the attachment of GPIs to proteins. The mechanisms and roles of the inositol acylation and deacylation have not been well clarified. Herein, we report derivation of human and Chinese hamster mutant cells defective in inositol acylation and the gene responsible, PIG-W. The surface expressions of GPI-anchored proteins on these mutant cells were greatly diminished, indicating the critical role of inositol acylation. PIG-W encodes a 504-amino acid protein expressed in the endoplasmic reticulum. PIG-W is most likely inositol acyltransferase itself because the tagged PIG-W affinity purified from transfected human cells had inositol acyltransferase activity and because both mutant cells were complemented with PIG-W homologs of *Saccharomyces cerevisiae* and *Schizosaccharomyces pombe*. The inositol acylation is not essential for the subsequent mannosylation, indicating that glucosaminyl phosphatidylinositol can flip from the cytoplasmic side to the luminal side of the endoplasmic reticulum.

INTRODUCTION

Glycosylphosphatidylinositol (GPI), a complex glycolipid, acts as a membrane anchor for many cell surface proteins (Tiede *et al.*, 1999; McConville and Menon, 2000; Ikezawa, 2002). In mammalian cells, >100 different proteins are GPI anchored, including cell surface enzymes, receptors, and adhesion molecules. GPI-anchored proteins are enriched in "lipid rafts," allowing their functional linkages to signal transduction systems (Simons and Toomre, 2000). GPI-anchored proteins are abundant in protozoan parasites, such as *Trypanosoma* and *Plasmodium* species (Ferguson, 1999; Gowda and Davidson, 1999). Biosynthesis of GPI is essential for the bloodstream form of *T. brucei* (Nagamune *et al.*, 2000), which causes sleeping sickness and has been validated as a therapeutic target (Ferguson, 2000). For the development of trypanosome-specific GPI biosynthesis inhibitors, it is im-

portant to clarify the different characteristics of human and trypanosomal pathways.

GPI is synthesized in the endoplasmic reticulum (ER) from phosphatidylinositol (PI) through multiple steps and is attached to proteins bearing the C-terminal GPI attachment signal sequence (Ferguson, 1999; Kinoshita and Inoue, 2000). The first two precursors of GPI are synthesized by transfer of N-acetylglucosamine (GlcNAc) to the inositol of PI to generate N-acetylglucosaminyl-PI (GlcNAc-PI), followed by deN-acetylation to generate glucosaminyl-PI (GlcN-PI). At the third step, the mammalian and *T. brucei* GPI biosynthetic pathways become different. In mammalian cells, the inositol acylation, mostly palmitoylation, occurs at GlcN-PI to generate GlcN-acylPI, and this precedes the first mannosylation that generates Man-GlcN-acylPI (Urakaze *et al.*, 1992; Stevens and Zhang, 1994; Doerrler *et al.*, 1996; Smith *et al.*, 1997). Once added, the inositol-linked acyl chain remains until the mature GPI anchor is attached to the protein. Usually, the acyl chain is removed from the GPI-anchored proteins by an inositol deacylase (Chen *et al.*, 1998). In contrast, inositol acylation occurs only after the first mannosylation in *T. brucei* (that is, the inositol acylation occurs at Man-GlcN-PI to generate Man-GlcN-acylPI) (Guther and Ferguson, 1995; Smith *et al.*, 1996b). The inositol acylation is required for the addition of the bridging phosphoethanolamine (P-EtN) to the third mannose (Guther and Ferguson, 1995).

The two initial reactions that generate GlcN-PI occur on the cytoplasmic side of the ER membrane (Vidugiriene and

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Abbreviations used: DAF, decay accelerating factor; GlcN, glucosamine; GlcNAc, N-acetylglucosamine; GlcN-PI(C8), glucosamine-PI bearing dioctanoyl-PI; GPI, glycosylphosphatidylinositol; GPI-PLD, glycosylphosphatidylinositol-specific phospholipase D; MTI, first mannosyltransferase; P-EtN, phosphoethanolamine; PI, phosphatidylinositol; PI-PLC, phosphatidylinositol-specific phospholipase C.

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Menon, 1993; Watanabe *et al.*, 1996; Nakamura *et al.*, 1997), whereas PIG-M, which transfers the first mannose to GlcN-acylPI, functions on the luminal side of the ER (Maeda *et al.*, 2001). Therefore, flipping should occur before the first mannosylation. Mutant cells defective in inositol acylation would be useful to determine whether inositol acylation is essential for flipping in mammalian cells and to further characterize the substrate specificities of the enzymes downstream in the biosynthetic pathway. Herein, we report the molecular cloning of rat PIG-W critical for inositol acylation and present evidence that inositol acylation is not essential for flipping of GPI but is important for the expression of GPI-anchored proteins on the cell surface.

MATERIALS AND METHODS

Cells and Culture

Human T lymphoma Molt4 cells were cultured in RPMI 1640 medium supplemented with 10% fetal calf serum (FCS). Chinese hamster ovary (CHO) cells were cultured in Ham's F-12 medium supplemented with 10% FCS. Human B-lymphoblastoid JY25 cells (Hollander *et al.*, 1988) and Thy-1-negative mutant cells of complementation class E derived from mouse lymphoma BW5147 (Hyman, 1988) were cultured in DMEM supplemented with 10% FCS. A 1D10 mutant cell was established by sorting CD59 negative cells from Molt4 cells (American Type Culture Collection, Manassas, VA), followed by limiting dilution. To establish CHOPA10.14 mutant cells, we screened for cells resistant to aerolysin, a bacterial toxin that binds to GPI-anchored proteins (Buckley, 1999), as described previously (Hong *et al.*, 2002). Cells were stained for decay accelerating factor (DAF) and CD59 and analyzed in a FACScan cytometer (BD Biosciences, San Jose, CA) (Nakamura *et al.*, 1997).

Plasmids

Rat PIG-W cDNA was subcloned into pME18Sf+, a mammalian expression vector, and pME-neo, pME18Sf+ bearing neo resistance gene, and the resulting plasmids were termed pME-rPIG-W and pME-neo-rPIG-W, respectively. To add FLAG tag to the N terminus of rat PIG-W, we replaced PIG-A cDNA of pME-FLAG-hPIG-A (Maeda *et al.*, 1998) with rat PIG-W cDNA, followed by subcloning a *Xba*I-*Nsi*I fragment bearing FLAG-PIG-W into pME-neo. To add FLAG tag to the C termini of rat PIG-W, we replaced *GPI*8 cDNA of pME-hGPI8-FLAG with rat PIG-W, followed by subcloning a *Xba*I-*Nsi*I fragment bearing PIG-W-FLAG into pME-neo. These plasmids were termed pME-neo-FLAG-rPIG-W and pME-neo-rPIG-W-FLAG, respectively. To generate pMEEB-FLAG-rPIG-W-GST and pMEEB-FLAG-rPIG-W-FLAG-HAT, we replaced *Xba*I-*Mlu*I fragments bearing *GPI*8 cDNA of pME-hGPI8-CST (Ohishi *et al.*, 2000) and pME-hGPI8-FLAG-HAT (Watanabe *et al.*, 2000), respectively, with *Xba*I-*Xba*I fragment of pME-neo-FLAG-rPIG-W and *Xba*I-*Mlu*I fragment of pME-neo-rPIG-W-FLAG, respectively. We then prepared *Xba*I-*Asp*718 fragments bearing FLAG-rPIG-W-GST and FLAG-rPIG-W-FLAG-HAT and cloned them into *Xba*I-and *Asp*718-cut pME-EBO, respectively. pMEEB-hPIG-L has been previously reported (Nakamura *et al.*, 1997).

Cloning PIG-W cDNA

Molt4 1D10 cells (10^8) were mixed with 300 μ g each of the plasmids of the rat C6 glioma cDNA library bearing simian virus 40 (SV40) origin of replication (Nakamura *et al.*, 1997) and pBSII-SV40T (ori-) for expression of SV40 large T protein in 4 ml of HEPES-buffered saline and were electroporated in 10 cuvettes at 250 V and 960 μ F by using a Gene Pulser (Bio-Rad, Hercules, CA). The next day, dead cells were removed by centrifugation through Ficoll-Paque (Pharmacia, Peapack, NJ). Two days after transfection, cells were stained with fluorescein isothiocyanate-labeled anti-CD59 antibody (BD Biosciences Pharmingen, San Diego, CA) and propidium iodide to gate out the dead cells, and 582 cells that restored CD59 expression were collected by a cell sorter (FACS-Vantage; BD Biosciences). Plasmid clones (10^3) were recovered from the cells. Pooled plasmids (90 μ g) were retransfected with 270 μ g of pBSII-SV40T (ori-) plasmid into 6×10^7 Molt4 1D10 cells in six cuvettes. After another cycle of cell sorting and recovery, 960 independent clones were analyzed and two positive clones were obtained.

Based on this rat sequence, we found two expressed sequence tag clones of human homolog (accession numbers BG196529 and BG741116). We also found two human genomic clones of chromosome 17 (accession numbers AC023133 and AC110594.2). We connected these sequences into a complete sequence of human PIG-W. We confirmed this sequence by amplification of the coding region from a human cDNA library by polymerase chain reaction and sequencing.

Analysis of GPI Biosynthesis

For labeling with mannose, 10^6 cells were incubated for 1 h in a medium containing 100 μ g/ml glucose and 10 μ g/ml tunicamycin (Sigma-Aldrich, St. Louis, MO) and then incubated in the same medium containing 40 μ Ci/ml d-[2-³H]mannose for 45 min (Hirosi *et al.*, 1992). Lipids were extracted and partitioned into *n*-butanol. To study the effect of YW3548/BE49385A, a terpenoid lactone (Sutterlin *et al.*, 1997), cells were incubated overnight in a medium containing 10 μ M BE49385A (a gift from Banyu Pharmaceutical Co., Tokyo, Japan), followed by labeling with mannose. For in vivo labeling with inositol, 10^6 cells were cultured in inositol-free DMEM supplemented with 10% dialyzed FCS and 20 μ Ci of myo-[2-³H]inositol for 1 d (Maeda *et al.*, 2001). Glycolipids were separated on Kiesel gel 60 (Merk, Darmstadt, Germany) and detected by a Fuji BAS1500 image analyzer (Fuji Film, Tokyo, Japan).

For analysis of the third step, we prepared membranes from cells preincubated with 5 μ g/ml tunicamycin for 2 h by hypotonic lysis and destruction with a Teflon homogenizer. The membranes obtained from Molt4 cells, its derivative cells, and CHOPA10.14 cells (10^7) were suspended in a buffer (100 mM Tris-HCl, pH 8/1 mM EDTA/protease inhibitors) and incubated for 10 min with 0.1 μ g of GlcN-PI (C8), GlcN-PI bearing dioctanoyl-PI (Doerfler *et al.*, 1996) (a gift from Dr. M.A. Lehrman; University of Texas Southwestern Medical Center, Dallas, TX) in 0.03% Triton X-100 and [³H]palmitoyl-CoA.

Enzyme Treatments of Labeled Lipids

Lipids were dissolved in 0.15 ml of a buffer containing either 100 mM Tris-HCl (pH 7.4), 0.1% Triton X-100, and 5 μ l (0.1 U) of *Bacillus thuringiensis* PI-specific phospholipase C (PI-PLC) (Toagosei, Tokyo, Japan); or 100 mM sodium acetate (pH 5.0), 1 mM ZnCl₂, 0.1% sodium taurodeoxycholate, and 15 μ l (1.7 U) of Jack bean α -mannosidase (Sigma-Aldrich); or 50 mM Tris-HCl (pH 7.4), 10 mM NaCl, 2.5 mM CaCl₂, 0.1% Triton X-100, and 15 μ l of human serum as a source of GPI-specific phospholipase D (GPI-PLD). After incubation for 16 h at 37°C, the lipids were extracted with *n*-butanol and analyzed by thin layer chromatography (TLC).

Purification of PIG-W Protein

JY25 cells (10^7) were transfected with 20 μ g of pMEEB-FLAG-rPIG-W-GST, pMEEB-FLAG-rPIG-W-FLAG-HAT, or pMEEB-hPIG-L-GST-FLAG by electroporation at 260 V and 960 μ F and cultured for 14 d with 400 μ g/ml hygromycin for selection. Proteins were affinity-purified from 10^8 cells as described previously (Watanabe *et al.*, 2000). The affinity-purified proteins were subjected to 4–20% gradient SDS-PAGE and were either silver stained or used for Western blotting. The proteins were also used for acyltransferase assay.

Acyltransferase Activity Assay With Liposomes Containing Purified PIG-W

A mixture of 3.9 mg of egg yolk phosphatidylcholine and 9.1 mg of phosphatidylethanolamine (Sigma-Aldrich) in 1 ml of chloroform was dried under N₂, dissolved in 6.5 ml of 100 mM Tris-HCl (pH 8.0) with 0.5 mM EDTA under sonication in a bath (UT-53N; Sharp, Osaka, Japan) for 5 min. Affinity-purified PIG-W or PIG-L proteins (equivalent to 2.5×10^7 cells) were agitated with 300 μ l of a buffer containing liposomes for 3 h at 4°C. A sample of 100 μ l of this liposome solution was incubated with 0.1 μ g of GlcN-PI (C8) and [³H]palmitoyl-CoA. After 10-min incubation at 37°C, lipids were extracted and analyzed by TLC.

Membrane Topology of PIG-W Protein

We generated CHO cells stably transfected with pME-neo-FLAG-rPIG-W or pME-neo-rPIG-W-FLAG. Cells were cultured on glass coverslips for 2 d and fixed in 4% paraformaldehyde for 15 min at room temperature. To selectively permeabilize the plasma membrane, cells were incubated in a buffer containing 5 μ g/ml digitonin. For complete permeabilization, 0.1% Triton X-100 was added to the blocking solution. Cells were then incubated with 2.5 μ g/ml rabbit anti-BiP antibody (ABR-Affinity BioReagents, Golden, CO), mouse anti-FLAG M2 antibody (Sigma-Aldrich), or rat anti-Hsc70 antibody (Stressgen Biotechnology, Victoria, BC, Canada) in 1% bovine serum albumin in phosphate-buffered saline (PBS) for 1 h. After three washes in PBS, cells were incubated with Texas Red-conjugated goat anti-rabbit IgG (Jackson Laboratories, Bar Harbor, ME), Alexa 488-conjugated anti-mouse IgG (Molecular Probes, Eugene, OR) or rhodamine-conjugated goat anti-rat IgG (Chemicon, Temecula, CA) in 1% bovine serum albumin in PBS for 1 h. Slides were mounted in mowiol and studied under a confocal microscope (Carl Zeiss, Jena, Germany).

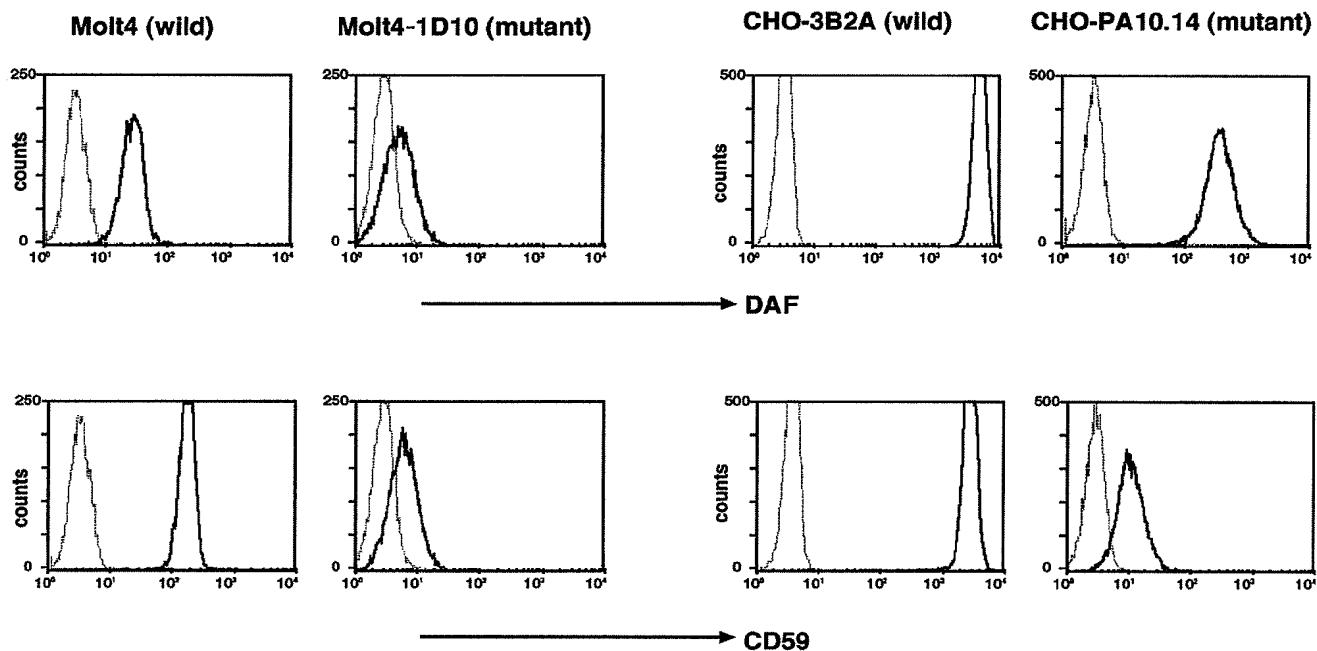


Figure 1. Defective surface expression of GPI-anchored proteins on Molt4-1D10 and CHOPA10.14 cells. Wild-type Molt4, mutant Molt4-1D10, wild-type CHO-3B2A, and mutant CHOPA-10.14 cells were stained for DAF (top) and CD59 (bottom) and analyzed in a FACScan. Solid lines, specific monoclonal antibodies; dotted lines, isotype-matched control antibodies.

RESULTS

Molt4-1D10 and CHOPA10.14 Cells Are Defective in the Expression of GPI-anchored Proteins Due to the Defective Inositol Acylation

We established Molt4-1D10 mutant from Molt4 cells and CHOPA10.14 mutant from CHO cells. Both mutants were partially defective in their surface expression of GPI-anchored proteins, DAF, and CD59 (Figure 1). PI-PLC removed the partially expressed DAF and CD59 from both cell lines, indicating that they were GPI anchored (our unpublished data).

To determine the molecular basis of the defective expression of GPI-anchored proteins on these mutant cells, we analyzed the biosynthesis of the GPI-anchor by metabolic labeling cells with mannose. In contrast to wild-type Molt4 cells (Figure 2A, lane 2), the Molt4-1D10 mutant accumulated abnormal mannosides, termed *a, *b, *c, *d, *e, and *f (lane 3). Those mannosides did not align with the mannosides in class K mutant cells (lane 1), which accumulate various intermediate and mature GPs, such as H5 to H8, due to a defect in the transfer of GPI to proteins (Yu *et al.*, 1997). CHOPA10.14 cells also accumulated a similar set of abnormal mannosides (Figure 2B, lanes 2 and 4).

All of those abnormal mannosides were sensitive to GPI-PLD (Figure 2A, lanes 6 and 7; B, lanes 1 and 2), like the GPI intermediates accumulated in class K mutant cells (Figure 2A, lanes 4 and 5), confirming that *a to *f were GPI species. In contrast to the GPI intermediates accumulated in class K mutant cells that were resistant to PI-PLC due to the inositol acylation (Figure 2A, lanes 8 and 9), all of *a to *f species were sensitive to PI-PLC (Figure 2A, lanes 10 and 11; B, lanes 3 and 4). These results indicated that the two mutants are defective in their inositol acylation and hence inositol acyl-

ation is important for the surface expression of GPI-anchored proteins.

Analysis of Mannolipids Accumulated in Cells Defective in Inositol Acylation

To determine steps for which the acyl chain on inositol is critical, we characterized mannosides accumulated in the 1D10 mutant. We first tested whether some of the mannosides have a P-EtN side chain on the first mannose. For this, we used YW3548/BE49386A, which inhibits the addition of P-EtN to the first mannose (Hong *et al.*, 1999). We pretreated mutant cells with this drug for 24 h before metabolic labeling with mannose (Figure 3A). Mannolipids *a and *b remained, whereas *c, *d, *e, and *f disappeared (lanes 1 versus 2), indicating that these four had the P-EtN side chain. Therefore, the inositol acylation is not required for the addition of the P-EtN side chain.

We then treated the mannosides with Jack bean α -mannosidase, which removes nonmodified mannoses (Figure 3B). All *a to *f mannosides were sensitive to the α -mannosidase, and two products, *g and *h, were formed (lane 1 versus 2). Because *a and *b were more hydrophobic than *g and *h, *g and *h were not products of *a and *b. Mannolipids *a and *b must have lost all mannoses upon the α -mannosidase treatment, suggesting that they were GlcN-PI-bearing nonmodified mannoses only. Mannolipids *a and *b probably have two and three mannoses, respectively, because CHOPA10.14 cells clearly had one more mannoside that was more hydrophobic than *a (Figure 2B, lanes 2 and 4, asterisk), most likely GlcN-PI bearing one mannose (Man-GlcN-PI).

To understand relationships between *g, and *c and *d, we isolated *c (Figure 3B, lane 4) and *d (lane 6) from the

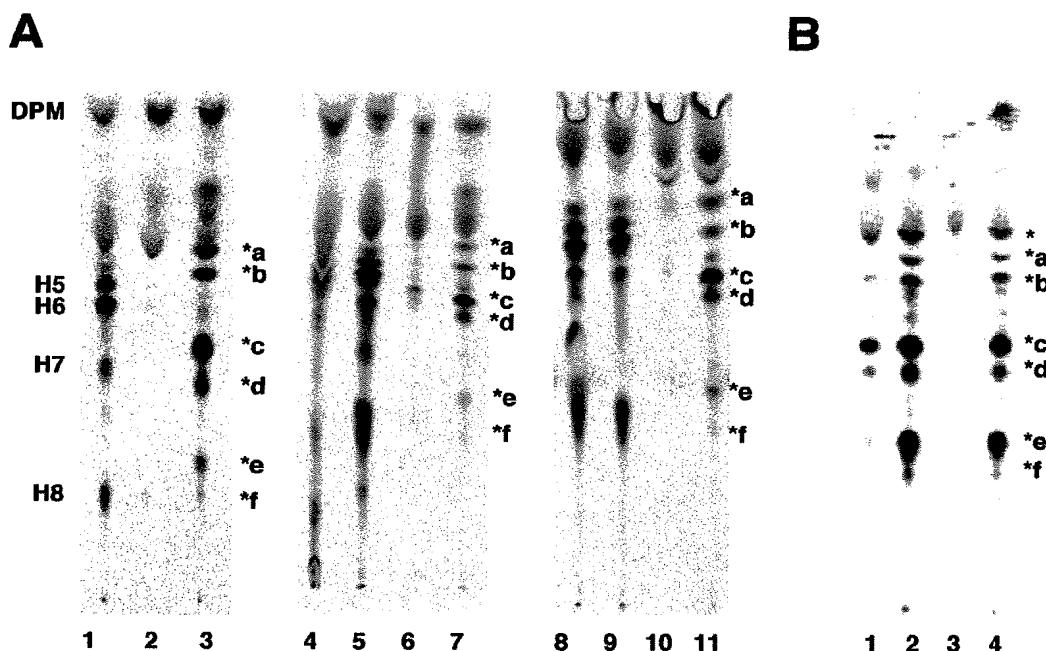


Figure 2. Accumulation of abnormal, PI-PLC-sensitive GPI species in Molt4-1D10 (A) and CHOPA10.14 (B) cells. (A) Class K mutant K562 (lane 1), wild-type Molt4 (lane 2), and 1D10 mutant (lane 3) cells were cultured with [³H]mannose in the presence of tunicamycin. Mannolipids were analyzed by TLC with a solvent system of chloroform/methanol/H₂O (10:10:3). DPM, dolichol-phosphate-mannose; H5, H6, H7, and H8 are according to Hirose *et al.*, 1992. *a-f, abnormal mannolipids accumulated in 1D10 cells. Labeled lipids of class K cells (lanes 4, 5, 8, and 9) and 1D10 cells (lanes 6, 7, 10, and 11) were treated with GPI-PLD (lanes 4 and 6), buffer for GPI-PLD (lanes 5 and 7), PI-PLC (lanes 8 and 10), or buffer for PI-PLC (lanes 9 and 11). Reextracted lipids were analyzed by TLC. (B) Mannolipids of CHOPA10.14 mutant cells prepared in a similar way to that described in A were treated with GPI-PLD (lane 1), buffer for GPI-PLD (lane 2), PI-PLC (lane 3), or buffer for PI-PLC (lane 4). *, abnormal mannolipid likely to be Man-GlcN-PI (see RESULTS).

TLC plate and treated them with the α -mannosidase. Mannolipid *g was derived from both *c (lane 3) and *d (lane 5). This result, taken together with the above-mentioned result that *c and *d have P-EtN on the first mannose (Figure 3A), suggested that *c was Man-(EtNP)Man-GlcN-PI, that *d was Man-Man-(EtNP)Man-GlcN-PI, and that the product *g was (EtNP)Man-GlcN-PI. Mannolipid *e became *h after α -mannosidase treatment, indicating that it had nonmodified mannose, most likely the third mannose. A big difference in the mobilities of *g and *h may be due to the presence of an additional P-EtN in *h. If this is true, *e may have the structures Man-(EtNP)Man-(EtNP)Man-GlcN-PI. Therefore, *b, *d, and *e probably had three mannoses, whereas there was no GPI species that had P-EtN on the third mannose, suggesting that inositol acylation may be critical for the addition of the bridging P-EtN that links GPI to proteins.

Finally, we tested whether a lack of the inositol acylation affects the efficiency of the first mannosylation. We metabolically labeled 1D10 cells and class E mutant cells (defective in synthesis of dolichol-phosphate-mannose; Tomita *et al.*, 1998) with inositol to assess levels of the first three GPI intermediates (Figure 3C). PI was seen mainly in wild-type Molt4 (lane 1), whereas GlcN-PI was accumulated in 1D10 mutant (lane 2). Transfection of the *PIG-W* cDNA normalized it (lane 3) (see the next section for cloning *PIG-W*). Class E cells accumulated GlcN-acylPI as expected (Urakaze *et al.*, 1992) (lane 4). These results indicate that inositol acylation is not essential for but accelerates the first mannosylation.

Cloning Rat *PIG-W* cDNA

We isolated a cDNA of the rat *PIG-W* gene that restored the surface expression of GPI-anchored proteins on Molt4-1D10 cells by means of expression cloning (Figure 4A, a and b). The same cDNA restored the surface expression of CD59 on CHOPA10.14 cells (c and d). Rat *PIG-W* cDNA also normalized GPI biosynthesis in the 1D10 mutant as revealed by metabolic labeling cells with mannose (Figure 4B). Transfection of a *PIG-W* cDNA (lane 3) but not an empty vector (lane 4) resulted in the disappearance of the accumulated manno-lipids seen in the 1D10 mutant (lane 2).

We then analyzed *PIG-W* mRNA in Molt4-1D10 cells by reverse transcription-polymerase chain reaction to see whether *PIG-W* is mutated. A cDNA of the same size was amplified from samples of wild-type and 1D10 mutant cells (our unpublished data). Direct sequencing of the band revealed a point mutation at nucleotide 281 that changed leucine to a stop codon, indicating that *PIG-W* is the defective gene in the 1D10 mutant.

Characteristics of *PIG-W* Protein

An open reading frame of rat *PIG-W* cDNA consisted of 1506 base pairs encoding 502 amino acids (Figure 5A). Searching databases, we found a human *PIG-W* gene (an intron-less gene mapped to chromosome 17q) that encoded 504 amino acids with 77% identity to rat *PIG-W* (the DDBJ/EMBL/GenBank accession numbers of rat and human *PIG-W* cDNAs are AB097817 and AB097818, respectively).

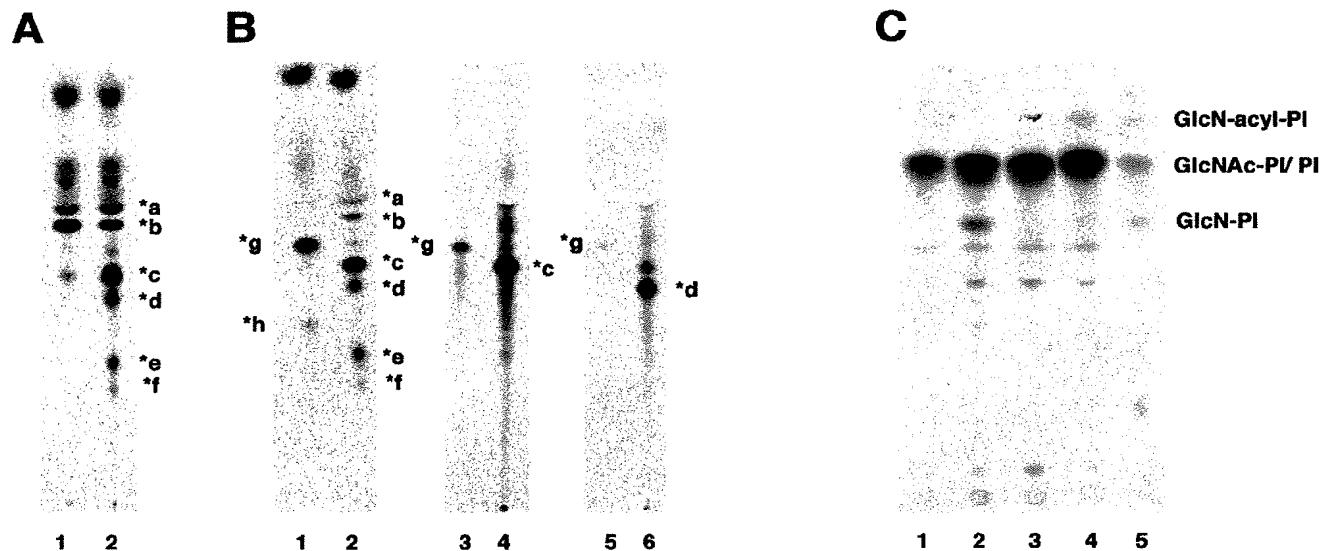


Figure 3. Characterization of mannosides accumulated in Molt4-1D10 (A and B) and analysis of early steps in GPI biosynthesis (C). (A) Molt4-1D10 cells cultured in the presence (lane 1) or absence (lane 2) of YW3548/BE49386A for 24 h were metabolically labeled with [³H]mannose. The lipids were analyzed by TLC with a solvent system of chloroform/methanol/H₂O (10:10:3). (B) Molt4-1D10 cells were metabolically labeled with [³H]mannose. Lipids were treated with Jack bean α -mannosidase (lane 1) or buffer (lane 2), extracted again, and analyzed by TLC as described in A. Mannosides *c and *d were eluted from the TLC plate (lanes 4 and 6) and treated with the α -mannosidase and reanalyzed by TLC (lanes 3 and 5). (C) Wild-type Molt4 (lane 1), 1D10 mutant (lane 2), 1D10 transfected with rat PIG-W cDNA (lane 3), and class E mutant (lane 4) cells were cultured in a medium containing myo-[³H]inositol for 1 d. The radiolabeled lipids were analyzed by TLC with a solvent system of chloroform/methanol/1 M NH₄OH (10:10:3). Lane 5, standard GlcNAc-PI, GlcN-PI, and GlcN-acyl-PI generated by incubating membranes of wild-type Molt4 with UDP-[6-³H]GlcNAc. PI and GlcNAc-PI are overlapping.

We also found yeast PIG-W homologs. *Saccharomyces cerevisiae* PIG-W (YJL091C) consists of 498 amino acids and has 28% amino acid identity to human PIG-W. *Schizosaccharomyces pombe* PIG-W (accession no. CAB59690) consists of 459 amino acids with 28% identity to human PIG-W (Figure 5A). Human PIG-W and its homologs had no significant sequence homology to other proteins, although they did share six well conserved regions (broken underlines in Figure 5A). The *S. cerevisiae* and *S. pombe* PIG-W homologs restored the surface expression of CD59 on Molt4-1D10 and CHOPA10.14 cells after transfection, indicating that they are functional PIG-W homologs (our unpublished data).

The hydropathic profile of human PIG-W shows multiple hydrophobic regions (Figure 5B). Analysis with the TMpred program (Smith *et al.*, 1996a) suggested the presence of 13 transmembrane domains in both human (Figure 5B, numbered bars) and *S. pombe* PIG-W (Figure 5A, numbered underlines). Based on comparison of locations of predicted transmembrane domains and the regions conserved among PIG-W homologs, presumably functionally important regions, we speculated that all the conserved regions are on the same side of the membrane as the N terminus (see below for the membrane orientation of the N terminus).

We then analyzed subcellular localization of PIG-W protein. We transfected a cDNA of glutathione S-transferase (GST)-tagged PIG-W into JY25 cells and fractionated nucleic-free lysates into the endoplasmic reticulum (ER), the Golgi apparatus, and plasma membranes by sucrose density gradient centrifugation (Figure 6A). Western blotting with anti-GST antibody revealed PIG-W mainly in fractions 4 and 5, corresponding to the ER (Figure 6B) (see the next section for an explanation for the higher molecular weight bands). Frac-

tions 1 and 2 containing the plasma membranes and the Golgi apparatus had none and only a small amount of PIG-W, respectively, indicating that PIG-W is an ER protein. There is no known ER localization signal sequence in PIG-W (Figure 5A).

Because PIG-W resides in the ER, the site of GPI biosynthesis, and because both *S. cerevisiae* and *S. pombe* PIG-W homologs were functional in human and hamster mutant cells, it is very likely that PIG-W is directly involved in inositol acylation.

PIG-W Is Most Likely the Acyltransferase

To set up an assay for inositol acyltransferase, we used a synthetic substrate GlcN-PI(C8). It was reported that GlcN-PI(C8) acted as a substrate of acyltransferase present in the membrane of mammalian cells, receiving the palmitoyl chain from [³H]palmitoyl-CoA (Doerrler *et al.*, 1996). Microsomal membranes of wild-type CHO (Figure 7, lane 1), but not the CHOPA10.14 mutant (lane 2), generated a specific spot (asterisk) in the presence but not absence (lane 3) of GlcN-PI(C8). Wild-type Molt4 (lane 4) inefficiently generated a similar spot (lane 4), whereas Molt4-1D10 transfected with PIG-W efficiently generated it in the presence (lane 5) but not absence (lane 6) of GlcN-PI(C8). An extra spot formed when acyltransferase activity was very high (double asterisks in lane 5). Nontransfected 1D10 did not make the specific spot (lane 7), indicating that this specific spot was GlcN-acyl-PI(C8). Therefore, a combination of GlcN-PI(C8) and [³H]palmitoyl-CoA should be useful to assay inositol acyltransferase.

To determine whether PIG-W is acyltransferase, we affinity purified PIG-W from human JY25 cells expressing rat

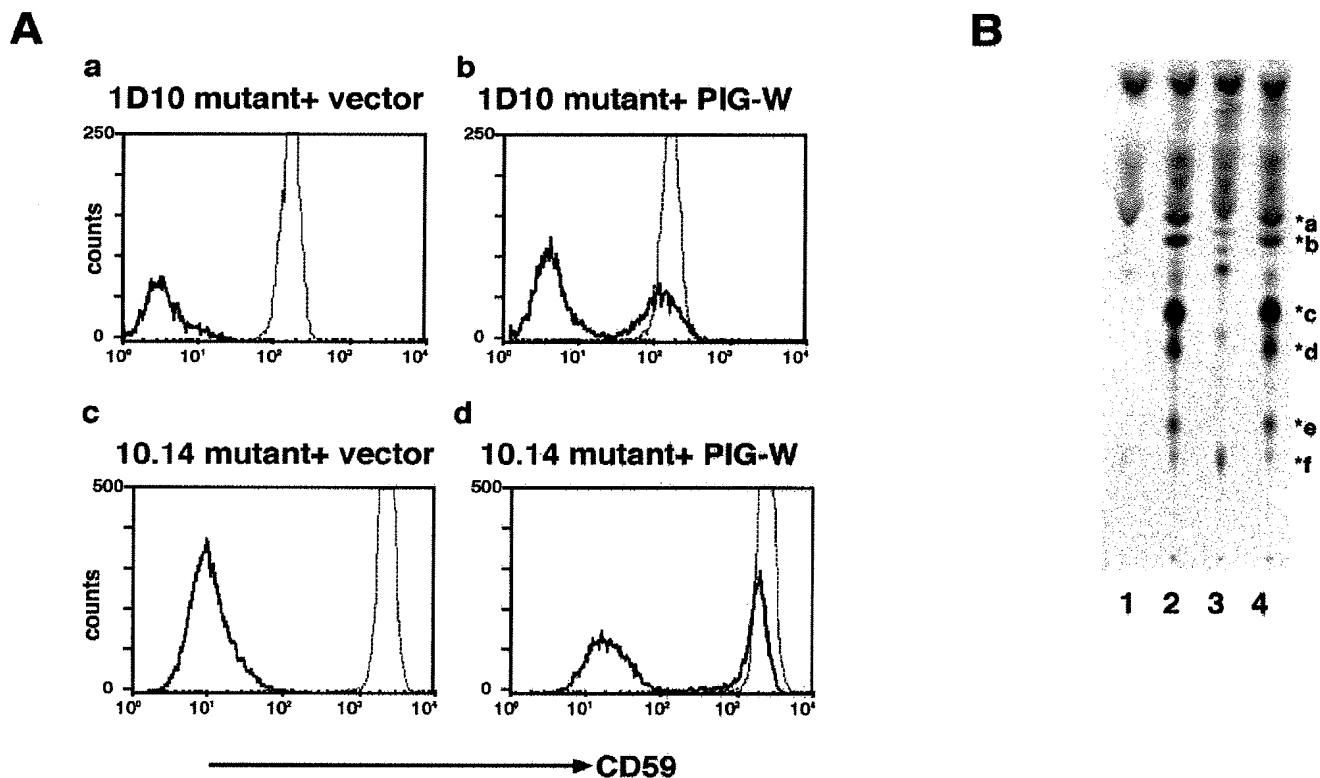


Figure 4. Molt4-1D10 and CHOPA10.14 cells are defective in PIG-W. (A) Rat PIG-W cDNA restored the surface expression of CD59 on Molt4-1D10 and CHOPA10.14 cells. 1D10 mutant transfected with an empty vector (a) or PIG-W (b) and CHOPA10.14 mutant transfected with an empty vector (c) or PIG-W (d) were stained 2 d after transfection. Solid lines, transfectants; dotted lines, wild-type cells. (B) Normalized GPI biosynthesis in 1D10 cells transfected with PIG-W cDNA. Wild-type Molt4 (lane 1), 1D10 mutant (lane 2), 1D10 transfected with rat PIG-W cDNA (lane 3), and 1D10 transfected with an empty vector (lane 4) were labeled with [³H]mannose and lipids analyzed by TLC.

PIG-W tagged with FLAG and HAT, or with FLAG and GST. As a control, PIG-L bearing FLAG and GST tags was similarly affinity purified. Both PIG-W preparations showed four silver-stained bands after SDS-PAGE under reducing conditions (Figure 8A, lanes 1 and 2). Bands of FLAG- and HAT-tagged PIG-W corresponded to 45, 90, 140–150, and 200 kDa (lane 1). All of them reacted with anti-FLAG antibodies on Western blotting (lane 4). The FLAG- and GST-tagged PIG-W showed four bands at 70, 150, 250, and >300 kDa (lane 2), all of which reacted with anti-FLAG antibodies (lane 5). The size difference between the two PIG-W preparations corresponded to the size difference between HAT (2 kDa) and GST (25 kDa). It is very likely that the four bands represented mono-, di-, tri-, and tetrameric PIG-W, respectively. The affinity-purified PIG-L showed a single band of an expected size (lanes 3 and 6).

We then measured the acyltransferase activity of the affinity-purified PIG-W by using GlcN-PI(C8) and [³H]palmitoyl-CoA. The specific product (asterisk) occurred after incubation with the FLAG- and GST-tagged PIG-W in the presence of GlcN-PI(C8) (Figure 8B, lane 1). This spot did not occur after incubation with PIG-L (lane 2) nor with the PIG-W in the absence of GlcN-PI(C8) (lane 3). To confirm that the product was in fact GlcN-acylPI(C8), the spot was extracted from the TLC plate and treated with GPI-PLC and

PI-PLC. After overnight incubation with buffer alone, the sample gave two extraradioactive spots (double and triple asterisks) (lanes 5 and 7), suggesting migration of the palmitoyl group during incubation (Costello and Orlean, 1992; Doerrler *et al.*, 1996). The one with double asterisks had mobility similar to that of the second spot seen with membrane-bound inositol acyltransferase (Figure 7, lane 5, double asterisks). The product was sensitive to GPI-PLD having generated a set of much less hydrophobic species (Figure 8B, lane 4). This is consistent with the prediction that GlcN-acylPI(C8) would yield radioactive GlcN-acyl-inositol. The reason for the generation of a cluster of three spots is unclear but may be related to migration of the acyl chain. The product was resistant to PI-PLC as expected (lane 6). Therefore, affinity-purified PIG-W had acyltransferase activity, suggesting that PIG-W is most likely an acyltransferase itself.

Luminal Orientation of the N and Cytoplasmic Orientation of the C Termini of PIG-W

We then determined the membrane orientation of the N and C termini of PIG-W. We generated rat PIG-W tagged with FLAG epitope at either the N or the C termini, confirmed that these tagged PIG-Ws are functional as determined by complementation of CHOPA10.14 cells, and then transfected

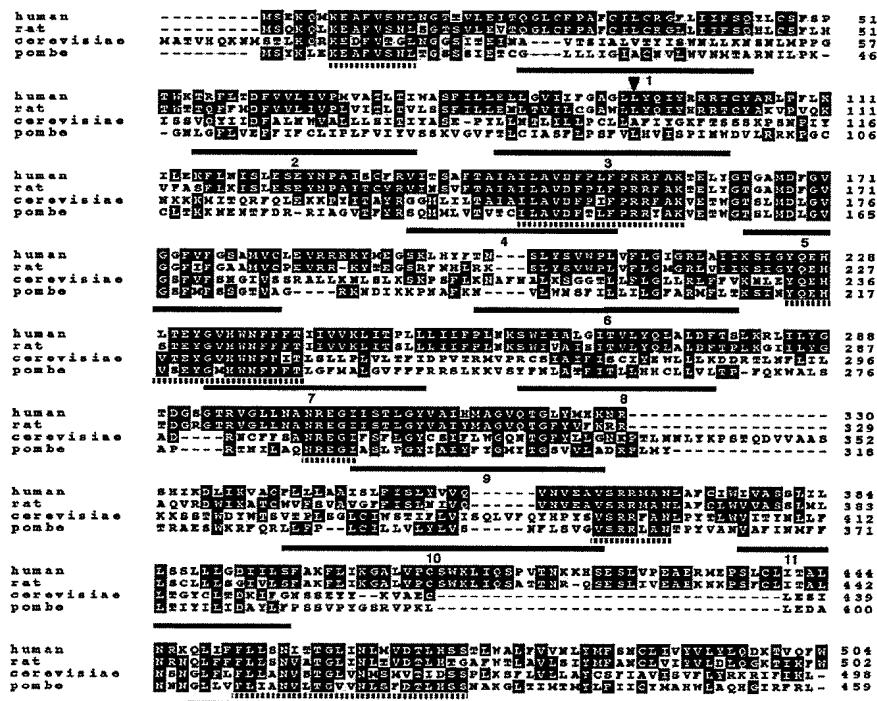
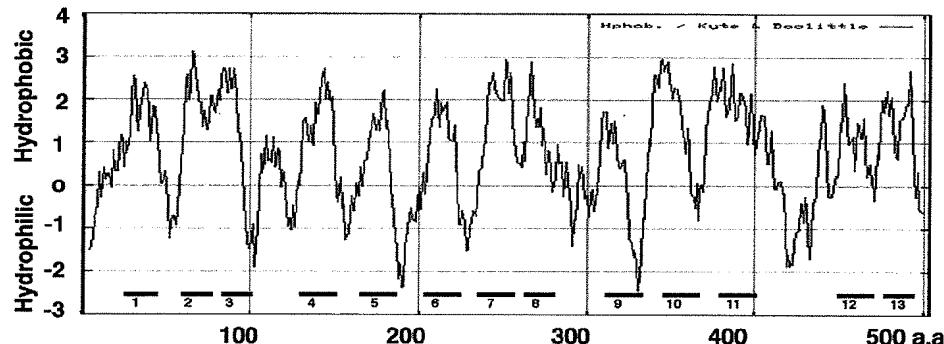
A**B**

Figure 5. (A) Alignment of amino acid sequences of PIG-W homologs of human, rat, *S. cerevisiae*, and *S. pombe*. Black and gray boxes indicate identical and conserved amino acids, respectively. Six regions highly conserved in four species are indicated by broken underlines. Positions of 13 predicted transmembrane domains of *S. pombe* PIG-W are indicated by numbered underlines. An arrowhead indicates leucine changed to a stop codon in 1D10 mutant. (B) Hydropathic profile of human PIG-W. Hydrophobicity was calculated with a window length of 17 (Kyte and Doolittle, 1982). Positions of 13 predicted transmembrane domains are indicated by numbered thick bars.

them into CHO cells (Figure 9). After permeabilization of both the plasma and ER membranes by Triton X-100 or after the selective permeabilization of only the plasma membrane by digitonin, we stained cells for a FLAG tag and an authentic ER luminal protein, BiP. We also stained an authentic

cytosolic protein Hsc70. The FLAG at the C termini of PIG-W was accessed by an antibody after permeabilization of just the plasma membrane (Figure 9e). Cells with the FLAG at the N terminus were stained only after permeabilization of the ER membrane (h versus k), similarly to BiP (g

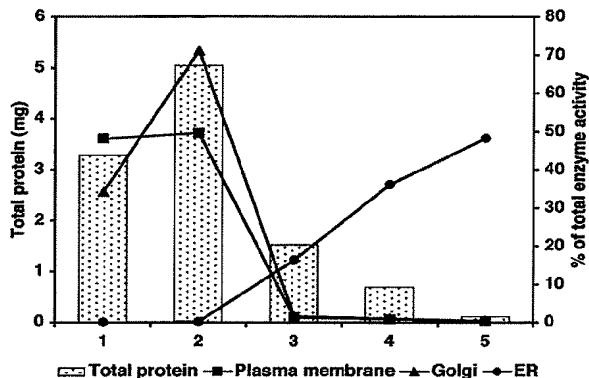
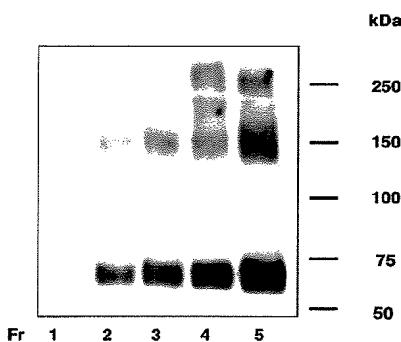
A**B**

Figure 6. ER localization of PIG-W. JY25 cells expressing GST-tagged PIG-W were hypotonically lysed and the postnuclear supernatant fractionated by sucrose density gradient centrifugation. Fractions 1–5 were characterized by protein content (bars) and organelle marker enzymes (alkaline phosphodiesterase I for the plasma membrane, α -mannosidase II for the Golgi apparatus, and dolichol-phosphate-mannose synthase for the ER) (A) and by Western blotting with anti-GST antibody (B). The enzyme activities in fractions are shown as percentages of the total.

versus j). Therefore, the N terminus was luminal and C termini was cytoplasmically oriented.

Combining this result with the prediction that the regions conserved among species and the N terminus have the same membrane orientation (Figure 5B), we speculated that the conserved regions may face the luminal side; hence, the acyltransfer probably occurs on the luminal side.

DISCUSSION

PIG-W Accounts for Most, If Not All, of the Inositol Acylation Activity

We isolated GPI-anchor-deficient cell lines, Molt4-1D10 and CHOPA10.14, and demonstrated that they are defective in the transfer of the acyl chain to inositol. We obtained rat *PIG-W* cDNA that restored GPI biosynthesis in Molt4-1D10 and CHOPA10.14 cells. Molt4-1D10 cells had a nonsense

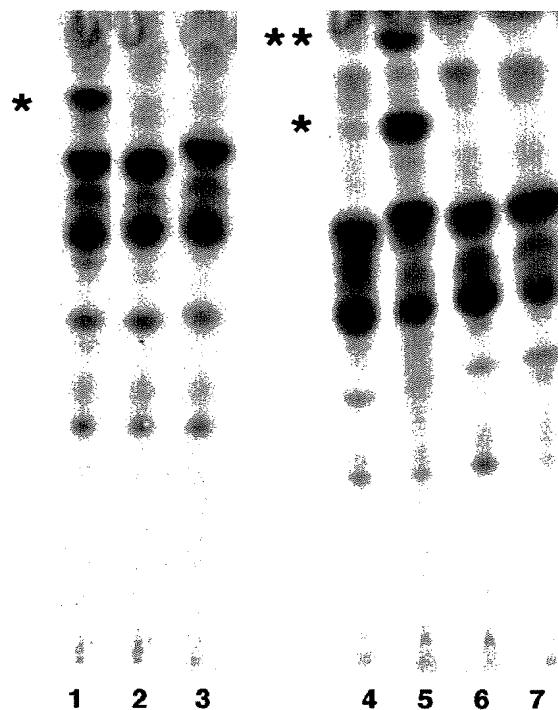


Figure 7. CHOPA10.14 and Molt4-1D10 are defective in inositol acylation of a synthetic substrate GlcN-PI(C8). The membranes of wild-type CHO (lanes 1 and 3) and CHOPA10.14 (lanes 2) cells were incubated with [³H]palmitoyl-CoA in the presence (lanes 1 and 2) or absence (lane 3) of GlcN-PI(C8). The membranes of wild-type Molt4 (lane 4), Molt4-1D10 transfected with PIG-W (lanes 5 and 6), and 1D10 mutant (lane 7) were incubated with [³H]palmitoyl-CoA in the presence (lanes 4, 5, and 7) or absence (lane 6) of GlcN-PI(C8). After 10-min-incubation, lipids were analyzed by TLC with a solvent system of chloroform/methanol/0.25% KCl (55:45:10). *, GlcN-acetylPI(C8); **, GlcN-PI(C8)-dependent extraspot of unknown structure.

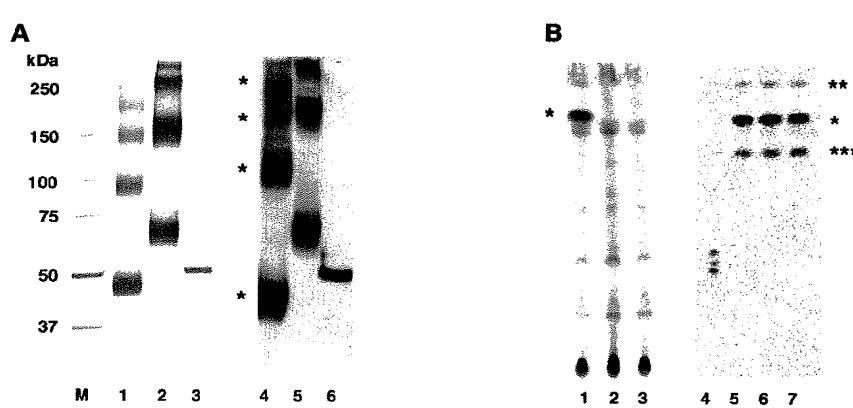
mutation at nucleotide 281 in the *PIG-W* gene. The epitope-tagged PIG-W, affinity purified from transfected human cells, had acyl-CoA-dependent inositol acyltransferase activity *in vitro*, indicating that Molt4-1D10 is defective in inositol acyltransferase.

Molt4-1D10 and CHOPA10.14 cells had no detectable inositol acylated GPI precursors. Therefore, PIG-W accounts for most, if not all, of the inositol acylation activity. Molt4-1D10 expressed only 20 and a few percent of normal levels of DAF and CD59, respectively, and CHOPA10.14 cells expressed only a few and <1% of DAF and CD59, respectively. These residual proteins were GPI anchored. It is unclear whether there is a second minor mechanism for the inositol acylation that may be responsible for the residual GPI anchoring (Stevens and Zhang, 1994) or whether noninositol-acylated GPI can, albeit inefficiently, be transferred to the proteins.

Inositol Acylation Is Not Essential for Flipping of GPI but May Be Critical for the Addition of the Bridging P-EtN to the Third Mannose

Both Molt4-1D10 and CHOPA10.14 cells accumulated a number of mannosylated noninositol-acylated GPI species.

Figure 8. The *PIG-W* gene encodes acyltransferase. (A) FLAG-PIG-W-FLAG-HAT (lanes 1 and 4), FLAG-PIG-W-GST (lanes 2 and 5), and PIG-L-GST-FLAG (lanes 3 and 6) were isolated by two-step affinity purification from the lysates of transfected JY25 cells. M, molecular size markers; lanes 1–3, silver staining; lanes 4–6, Western blotting with anti-FLAG antibody. (B) The purified FLAG-PIG-W-GST (lanes 1 and 3) and PIG-L-GST-FLAG (lane 2) proteins were incubated with [³H]palmitoyl-CoA in the presence (lanes 1 and 2) or absence (lane 3) of GlcN-PI(C8). After 10-min-incubation, lipids were analyzed by TLC with a solvent system of chloroform/methanol/0.25% KCl (55:45:10). GlcN-acylPI(C8) generated (asterisk) was treated overnight with GPI-PLD (lane 4), buffer for GPI-PLD (lane 5), PI-PLC (lane 6), or buffer for PI-PLC (lane 7), extracted again, and analyzed by TLC.



Considering that GPI should flip from the cytoplasmic side to the luminal side before the first mannosylation, this result indicated that inositol acylation is not essential for flipping. This is consistent with a report that the additions of three mannoses occurred in *T. brucei* even when inositol acylation was inhibited by phenylmethylsulfonyl fluoride (Guther *et al.*, 1994).

That the first mannosylation occurs without inositol acylation was a surprising finding, and contrary to our current concept that inositol acylation is a prerequisite for the addition of the first mannose in mammalian cells (Urakaze *et al.*, 1992; Smith *et al.*, 1997). It should still hold that inositol acylation precedes the first mannosylation in mammalian cells (Doerrler *et al.*, 1996). The first mannosylation by PIG-M (Maeda *et al.*, 2001) seemed to be more efficient with inositol acylation because GlcN-PI was accumulated in Molt4–1D10 cells and this accumulation disappeared after transfection of *PIG-W* cDNA.

Mannolipids *b, *d, and *e seem to have three mannoses, suggesting that inositol acylation is not essential for the second mannosyltransferase and the PIG-B-mediated third

mannosylation (Takahashi *et al.*, 1996). Mannolipids *c, *d, *e, and *f had the P-EtN side chain on the first mannose. Therefore, inositol acylation is not essential for the action of PIG-N, the enzyme that attaches this side chain (Hong *et al.*, 1999). It may be that inositol-acylated species are better substrates of these three enzymes because nonacylated species were accumulated.

On the contrary, we did not find GPI having P-EtN on the third mannose in these mutants. Therefore, inositol acylation may be critical for the addition of the bridging P-EtN to the third mannose, suggesting that the substrate specificity of the P-EtN transferase consisting of PIG-O and PIG-F (Hong *et al.*, 2000) is restricted to the inositol acylated intermediate.

Comparison of GPI Biosynthetic Pathways in Mammalian Cells and *T. brucei*

The first mannosyltransferase (MT1) of *T. brucei* mannosylates GlcN-PI efficiently, whereas mammalian MT1 prefers GlcN-acylPI (Doerrler *et al.*, 1996; Smith *et al.*, 1996b). This different substrate specificity of MT1 has been a rational

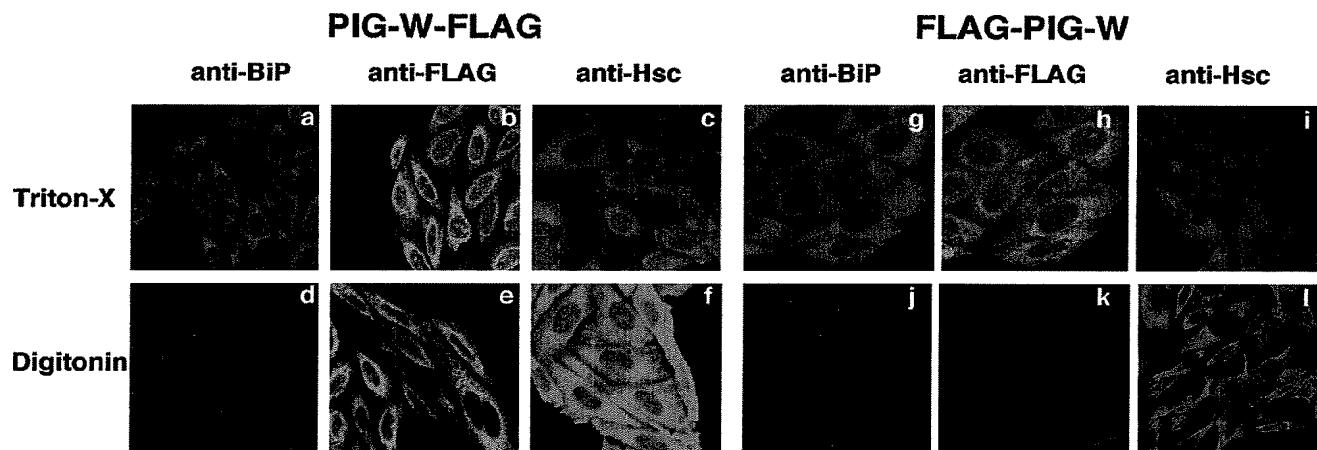


Figure 9. Membrane orientation of the N and C termini of PIG-W. Rat PIG-W tagged at the C termini (PIG-W-FLAG) (a–f) or the N terminus (FLAG-PIG-W) (g–l) were expressed in CHO cells. After permeabilization of both the plasma membrane and the ER membrane (a–c and g–i), or selective permeabilization of the plasma membrane (d–f and j–k), cells were stained for tagged PIG-W by anti-FLAG antibody (b, e, h, and k), for an endogenous ER luminal protein BiP (a, d, g, and j) and a cytosolic protein Hsc70 (c, f, i, and l).

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basis for the development of trypanosome-specific inhibitors. In fact, a substrate analog GlcN-(2-O-hexadecyl)PI, in which position 2 of inositol is modified by C16 alkyl instead of acyl chain, acts as an inhibitor of *T. brucei* MT1 but not mammalian MT1 (Smith *et al.*, 1997). The present study demonstrated that mammalian MT1 clearly uses GlcN-PI as a substrate and is not strictly dependent upon inositol acylation. Therefore, substrates of mammalian and *T. brucei* MT1 at least partially overlap, rather than clearly differ from each other. It is unknown whether *T. brucei* MT1 can mannosylate GlcN-acylPI partly because chemical synthesis of GlcN-acylPI is not practical due to acyl migration. PIG-M, an ER membrane protein with multiple transmembrane domains, is mammalian MT1 (Maeda *et al.*, 2001). *T. brucei* has a PIG-M homolog with 30% amino acid identity to human PIG-M (Maeda *et al.*, 2001).

Regarding substrate specificity of inositol acyltransferases, it was reported that the *T. brucei* enzyme acts on Man-GlcN-PI but not GlcN-PI (Guther and Ferguson, 1995), whereas mammalian inositol-acyltransferase acts on GlcN-PI (Urakaze *et al.*, 1992; Stevens and Zhang, 1994; Doerrler *et al.*, 1996). These specificities are complementary to the substrate specificity of MT1, accounting for the different orders of two reactions. It is unclear, however, whether *T. brucei* has any ability to generate GlcN-acylPI *in vivo*. Disruption or RNAi (Wang and Englund, 2001) of PIG-M in procyclic *T. brucei* need to be undertaken to determine whether indeed the first mannosylation is a prerequisite for inositol acylation. In bloodstream form *T. brucei*, GPI intermediates bearing one, two, and three mannoses are in equilibrium between the inositol-acylated and the-nonacylated forms (Guther and Ferguson, 1995). This is due to the actions of inositol acyltransferase and inositol deacetylase (Guther and Ferguson, 1995). All three mannosylated GlcN-PI may be substrates of *T. brucei* PIG-W, or alternatively there are more than one inositol acyltransferases with different substrate specificities. With respect to the inositol deacetylation, one enzyme has recently been characterized and the presence of yet another inositol deacetylase has been proposed (Guther *et al.*, 2001).

It was reported that inositol acylation in *T. brucei* is critical for the transfer of the P-EtN that bridges the GPI-anchor to the protein, i.e., the acyl chain on inositol is a requirement for ethanolaminephosphotransferase (Guther and Ferguson, 1995). The present study suggested that this may be also true in a mammalian system, where the transfer of the bridging P-EtN is mediated by a complex of PIG-O and PIG-F, in which PIG-F stabilizes catalytic PIG-O (Hong *et al.*, 2000). It is suggested that *T. brucei* may have a similar enzyme.

Membrane Orientation of Inositol Acylation

Comparison of amino acid sequences of mammalian PIG-W and PIG-W homologs of various organisms revealed six well conserved regions. Sites important for inositol acylation are very likely included in these regions. Based on the positions of predicted transmembrane domains and the luminal orientation of the N terminus of PIG-W, we predicted that all the well conserved regions face the luminal side of the ER. If this prediction is correct, inositol acylation may occur on the luminal side. Whereas the presence of acyl-CoA on the luminal side has not been demonstrated, there may be a mechanism for translocation of acyl-CoA across the ER membrane. An analysis of functional sites of PIG-W and determination of their membrane orientation are required to prove the luminal orientation of inositol acylation.

Note added to proof. Two articles describing PIG-W homologue of *Saccharomyces cerevisiae* have appeared recently: Tsukahara, K., Hata, K., Nakamoto, K., Sagane, K., Watanabe, N., Kuromitsu, J., Kai, J., Tsuchiya, M., Ohba, F., Jigamti, Y., Yoshimatsu, K., and Nagasu, T. (2003) Medical genetics approach towards identifying the molecular target of a navel inhibitor of fungal cell wall assembly. *Mol. Microbiol.* 48, 1029–1042; Umemura, M., Okamoto, M., Nakayama, K., Sagane, K., Tsukahara, K., Hata, K., and Jigami, Y. (2003) GWT1 gene is required for inositol acylation of glycosylphosphatidylinositol anchors in yeast. *J. Biol. Chem.* 278, 23639–23647.

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